## ISO/IEC 15504-10 Safety Extension, Yet Another Safety Standard?

### Giuseppe Lami, PhD

Consiglio Nazionale delle Ricerche Istituto di Scienza e Tecnologie dell'Informazione via Moruzzi, 1 - Pisa

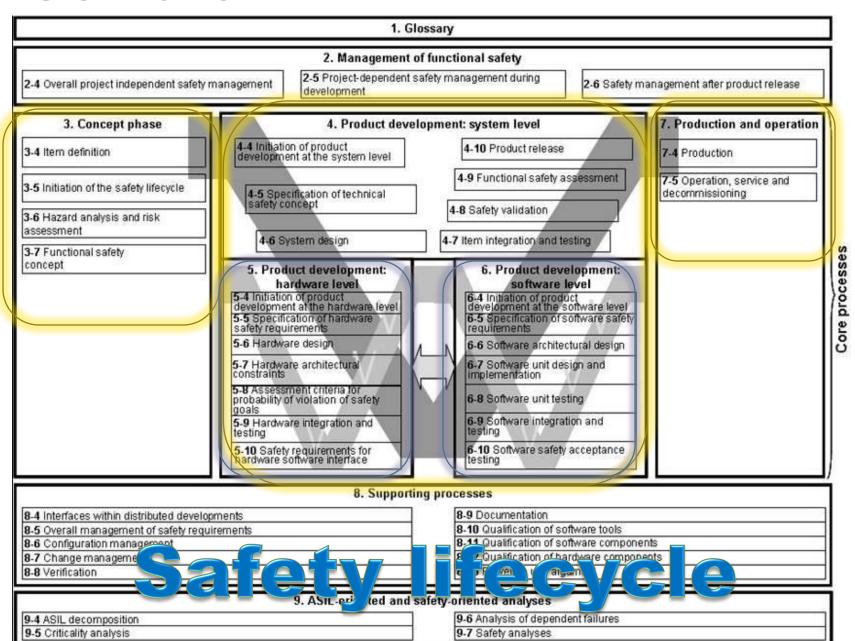
8° Workshop di Automotive SPIN Italia



### Talk outline

- Relationship between ISO 26262 & Automotive SPICE
- Overview of the last release of ISO/IEC 15504-10 Safety Extension (DTR ballot)
- How to integrate ISO 26262 & ISO/IEC 15504-10
- Conclusions

### ISO 26262



## How does Automotive SPICE support the Safety Lifecycle?

- The support provided by Automotive SPICE to the ISO 26262 Safety Lifecycle is related to:
  - Project Management
  - CM
  - QM
  - ENG.\*
  - Product release
  - Documentation
- From a producer point of view the goal is to understand the overlapping between.
   With a synergical perspective.

# Overlapping between Automotive SPICE processes & ISO 262626 requirements

Activities out of the Automotive SPICE scope

- •HW development
- •Hazard analysis & risk assessment
- Safety analysis
- Functional Safety concept
- Definition of technical safety concept
- •Safety management (overall, during dev., after SOP)
- •Safety qualification (tools, libreries, components)
- Safety validation (sw & sys)

•CM

- Project management
- Problem Resolution & Change management
- Documentation
- Quality management
- Software/system Req.Management
- Software Development
- Software/system V&V

Activities
within the
Automotive
SPICE
scope

• It has been argued that having the Automotive SPICE processes (HIS scope, as a minimum) at level 2 provides support for the fulfillemnt of the ISO 26262 requirements (see the *intersection area*).

•CM

- Project management
- Problem Resolution & Change management
- •Documentation
- Quality management
- Software/system Req. Management
- •Software Development
- •Software/system V&V

ISO 26262

[see E. Petry presentation @ 6W]



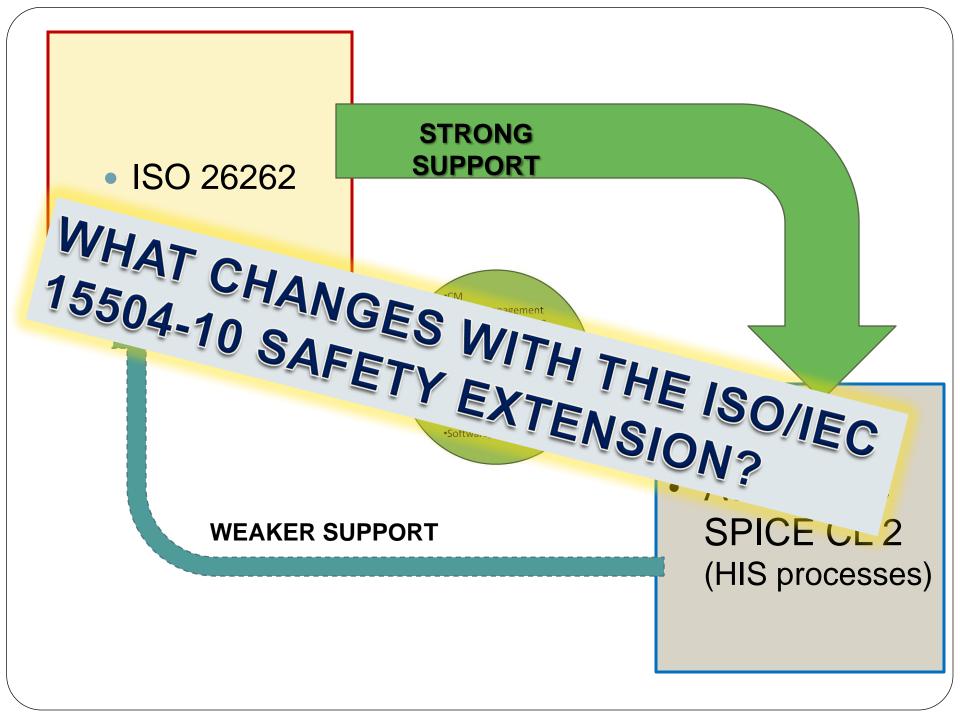
Automotive SPICE

## Scenario 1: ISO 26262 compliance

- Having achieved the ISO 26262 compliance for a certain project allow to infer the perfomance (and management) of processes (CM; PM; Problem resolution and change Man.; Documentation; QM; Requirements Man., Sw/System development, V&V) => achievement of Automotive SPICE capability level 1/2.
- It doesn't provides any guarantee of level 3 for processes
  - •CM
  - Project management
  - •Problem Resolution &
  - Change management
  - Documentation
  - Quality management
  - •Software/system Req. Management
  - Software Development
  - Software/system V&V

## Scenario 2: Automotive SPICE (HIS processes) level 2

- It supports the achievement of compliance of ISO 26262 requirements for
  - CM; PM; Problem resolution and change Man.;
     Documentation; QM; Requirements Man.
  - Uncertain support for SW/Sys development and V&V.
     (because the way processes are performed may be different from the way required by the ISO 26262)
- It doesn't provide any guarantee for ISO 26262 compliance



### ISO 26262 vs. ISO/IEC 15504-10

- The goal of ISO 26262 (61508) is to demonstrate the capability to develop certain products with acceptable risks
- To do that the organization is required to perform defined safety management, safety

engineering and safety

qualification activities

 These activities correspond to the processes belonging to the ISO/IEC 15504-10 Safety Extension

- •Hazard analysis & risk assess.
- Safety analysis
- •Functional Safety concept
- •Definition of technical safety concept
- •Safety management (overall, during dev., after SOP)
- •Safety qualification (tools,
- libreries, components)
- •Safety validation (sw & sys)

- •CM
- Project management
- •Problem Resolution & Change management
- Documentation
- Quality management
- •Software/system Req. Management
- •Software Development
- •Software/system V&V

## ISO/IEC 15504-10 Safety Extension

Safety Management process (SAF.1): The purpose is to ensure that products, services and life cycle processes meet safety objectives

#### Process outcomes:

- Safety principles and safety criteria are established.
- The scope of the safety activities for the project is defined.
- Safety activities are planned and implemented.
- Tasks and resources necessary to complete the safety activities are sized and estimated.
- Safety organization structure (responsibilities, roles, reporting channels, interfaces with other projects or OUs ...) is established.
- Safety activities are monitored, safety-related
   incidents are reported, analysed, and resolved.
- Agreement on safety policy and requirements for supplied products or services is achieved.
  - Supplier's safety activities are monitored.

#### **Base Practices:**

- SAF.1.BP.1: Define safety objectives and criteria.
- SAF.1.BP.2: Define Safety Life Cycle.
- SAF.1.BP.3: Perform safety planning.
- SAF.1.BP.4: Define safety activities integration.
- SAF.1.BP.5: Define skills requirements definition and allocate responsibility. SAF.1.BP.6: Implement planned safety activities.
- SAF.1.BP.7: Monitor the deployment of the safety activities.
- SAF.1.BP.8: Define and agree safety policy and safety requirements with suppliers.
- SAF.1.BP.9: Monitor the safety activities of the supplier.
- SAF.1.BP.10: Implement an escalation mechanism.

## ISO/IEC 15504-10 Safety Extension

Safety Engineering process (SAF.2): the purpose is to ensure that safety is adequately addressed throughout all stages of the engineering processes.

#### Process outcomes:

- Hazards related to product are identified and analysed.
- Hazard log is established and maintained.
- Safety demonstration for the product lifecycle is established and maintained.
- Safety requirements are defined.
- Safety integrity requirements are defined and allocated.
- Safety principles are applied to development processes.
- Impacts on safety of change requests are analysed.
- Product is validated against safety requirements.
- Independent evaluations are performed.

#### **Base Practices:**

- SAF.2.BP.1: Identify hazard sources and hazards.
- SAF.2.BP.2: Analyze hazards and risks.
- SAF.2.BP.3: Establish and maintain hazard log.
- SAF.2.BP.4: Establish and maintain safety demonstration.
- SAF.2.BP.5: Establish and maintain safety req.s.
- SAF.2.BP.6: Determine safety integrity requirements. NOTE: Safety integrity req. may be described i.e. as SIL.
- SAF.2.BP.7: Allocate safety requirements and safety integrity requirements.
- SAF.2.BP.8: Apply safety principles to achieve safety integrity requirements.
- SAF.2.BP.9: Perform safety impact analysis on changes.
- SAF.2.BP.10: Perform safety validations on product.
- SAF.2.BP.11: Perform independent assessments.

## ISO/IEC 15504-10 Safety Extension

**Safety Qualification process (SAF.3)**: the purpose is to assess the suitability of external resources when developing a safety-related software or system.

#### Process outcomes:

- Safety qualification strategy for external resources is developed.
- Safety qualification plan is developed and executed.
- Safety qualification documentation is written.
- Safety qualification report is produced.

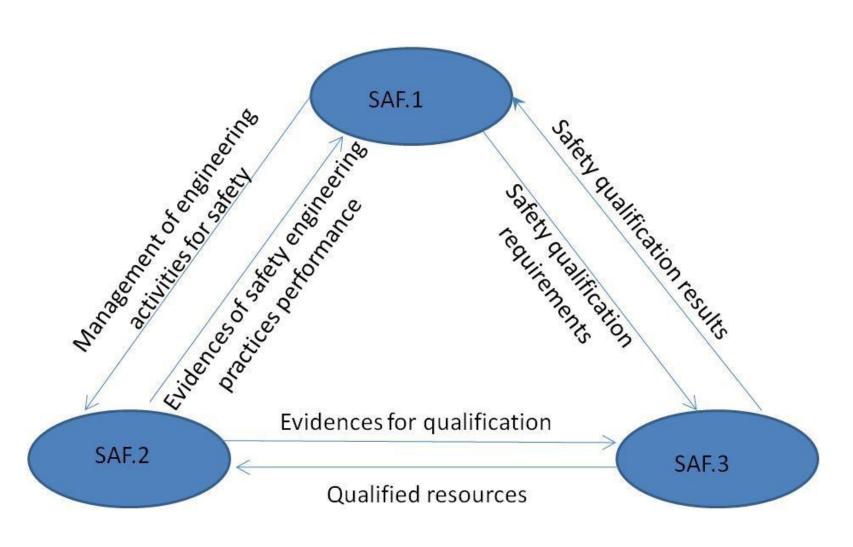
#### **Base Practices:**

- SAF.3.BP.1: Develop a safety qualification strategy.
- SAF.3.BP.2: Plan the safety qualification of external resources.

NOTE: Examples of external resources are as follows:

- core engineering tools automatic code generators, compilers and linkers;
- engineering support tools test, build and CM tools;
- management support tools documentation and project management tools.
- SAF.3.BP.3: Qualify the external resources.
- SAF.3.BP.4: Record the safety qualification results.
- SAF.3.BP.5: Maintain and update the safety qualification results.

## ISO/IEC 15504-10 Realtionship among processes



# ISO/IEC 15504-10 clause 5: Lifecycle Guidance

- It gives the assessors, for each process contained in 15504-5 and 15504-6, an indication of additional issues to be taken into account at assessment time.
- In Annex A WP characteristics are provided



### ISO 26262 vs. ISO/IEC 15504-10

- •Hazard analysis & risk assess.
- Safety analysis
- Functional Safety concept
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- •Safety management (overall, during dev., after SOP)
- •Safety qualification (tools, libreries, components)
- Safety validation (sw & sys)

- •CM
- Project management
- Problem Resolution & Change management
- Documentation
- Quality management
- •Software/system Req. Management
- Software Development
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## Realtionship between ISO 26262 & ISO/IEC 15504-10

## Scenario 1: ISO 26262 compliance

It is possible to infer the perfomance (and management) of processes into the HIS scope + those belonging to the ISO/IEC 15540-10 achievement of Automotive SPICE (HIS) CL1/2 + CL 1/2 of the ISO/IEC 15504-10 processes.

Why should I spend effort for increasing the capability of ISO/IEC 15504-10 processes?

Scenario 2: Automotive SPICE (HIS)+ ISO/IEC 15504-10 (CL 2)

- It supports the achievement of compliance of ISO 26262 requirements for:
  - all the activties but
  - SW/Sys development and V&V. activities (because the way process are prerformed may be different from the way required by the ISO 26262)
  - It doesn't provide any guarantee for ISO 26262 compliance

## Why increasing ISO/IEC 15504-10 processes capability?

- Capability is not (only) performance
- Higher capability means:
  - Efficient and repeatable development
  - Lower risk of missing project objectives
  - Identification of improvement areas
- Improving the capability of 15504-10 processes:
  - Isn't a short cut for ISO 26262 but
  - It is a way to improve safety-related processes
  - Is a managerial choice oreinted to process improvement
  - Is a way to extend the benefits of ten years of process improvement in automtive to safety-related activities

### Thanks

### Giuseppe Lami, PhD

## ISO/IEC 15504 & Automotive SPICE Principal Assessor

cert. N. Intacs-3961-1000-20254-03 cert. N. IntRSA 07/IntRSA-91004-I

Consiglio nazionale delle Ricerce Istituto di Scienza e Tecnologia dell'Informazione (ISTI) via Moruzzi,1 56124 Pisa (Italia)

phone: 0503153493

email: giuseppe.lami@isti.cnr.it